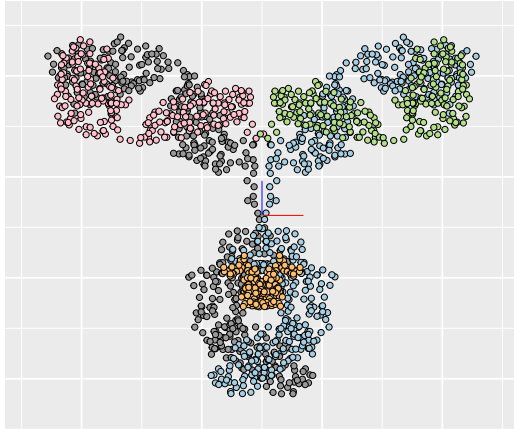


Populations

R.W. Oldford

Problem: Structure of human immunoglobulin G1 (IgG1)

Recall exploring how the geometry of the human immunoglobulin G1 molecule related to different variables associated with each “alpha” carbon.



E.g. here, colours are assigned to each carbon atom according to the value of its chainID variable.

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The data frame `igg1` (from the package `loon.data`) has 1556 rows, one for each alpha carbon. In the above notation, we can take

- ▶ row i to be the i th unit in \mathcal{P} , and
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- ▶ they can also be thought of as possible **keys** to identify identical units (e.g. as `linkingKeys` in `loon` plots).

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Notation: Populations will be distinguished from one another by using subscripts as in $\mathcal{P}_{\text{IgG1}}$.

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The data frame `igg1` also has 10 columns, each being a variable recording its value for every individual alpha carbon (unit) in the data frame.

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The same is true for the remaining variables in `igg1`: `recordType`, `name`, `residue`, `chainID`, `residueSequenceNum`, `residueName`, `group`. Each records the values of these **variates** for the units in our data set, namely $u \in \{u_1, u_2, \dots, u_{1556}\}$.

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- ▶ similarly, the other two coordinates of the 3D locations $y(u)$ and $z(u)$ (or simply y_u and z_u) are also *continuous* and *ratio scale* variates.

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- ▶ e.g. a variate such as $preference(u) \in \{"hate", "dislike", "neutral", "like", "love"\}$
- ▶ there are no strictly ordinal variates in the `igg1` data (though several, `residueSequenceNum`, `x`, `y`, and `z` can each be ordered)

Data: Realizations, observations, and variates

The first three rows of igg1 are

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## 2      ATOM  CA   VAL      H              2 -60.766 48.666 -15.351
## 3      ATOM  CA   GLN      H              3 -57.145 48.577 -16.631
##      residueName                      group
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## 2      Valine Non-polar (hydrophobic)
## 3    Glutamine      Polar (uncharged)
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The first three rows of `igg1` are

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## recordType name residue chainID residueSequenceNum      x      y      z
## 1      ATOM  CA    GLU      H              1 -62.259 45.262 -16.149
## 2      ATOM  CA    VAL      H              2 -60.766 48.666 -15.351
## 3      ATOM  CA    GLN      H              3 -57.145 48.577 -16.631
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N.B. Some people refer to this standard arrangement and interpretation as a **tidy data** representation.

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paste0(round(100 * prop), "%") # as a character string for printing
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Or, maybe, a two way table of counts for combinations of `chainID` and `group`

```
knitr::kable(with(igg1, table(chainID, group)))
```

	Acidic	Basic	Non-polar (hydrophobic)	Polar (uncharged)	Sugar
C	0	0	0	0	220
H	38	54	171	189	0
I	38	54	171	189	0
L	17	19	78	102	0
M	17	19	78	102	0

where some similarities and differences between chains are immediately apparent.

Chains H and I are “heavy”, L and M “light”, and C is a carbohydrate chain.

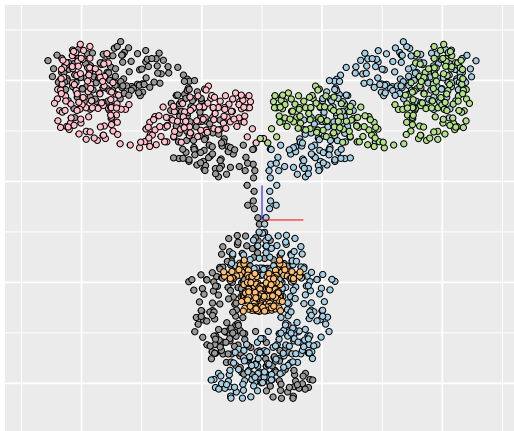
Population attributes – graphical

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For example, as already seen, the geometric locations shown in an interactive 3D scatterplot can be very informative (here coloured by chain ID):



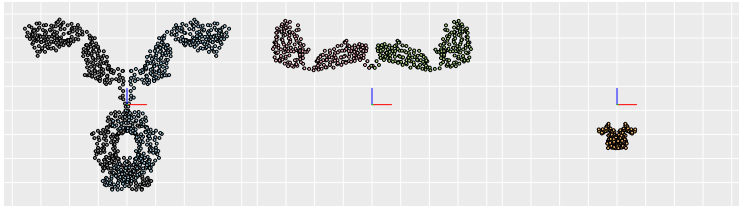
Population attributes – graphical

Interactive graphics, as in loon , make it very easy to construct informative graphical attributes by direct manipulation, as well as to save them for traditional publication:

```
heavyChain <- (igg1$chainID == "H") | (igg1$chainID == "I")
lightChain <- (igg1$chainID == "L") | (igg1$chainID == "M")
carbs <- (igg1$chainID == "C")
p3d["active"] <- heavyChain
p3d_heavy <- plot(p3d, draw = FALSE)
p3d["active"] <- lightChain
p3d_light <- plot(p3d, draw = FALSE)
p3d["active"] <- carbs
p3d_carbs <- plot(p3d, draw = FALSE)
# And plot these using grid graphics extra functionality
library(gridExtra)
# to arrange them in sequence
grid.arrange(p3d_heavy, p3d_light, p3d_carbs, nrow = 1)
```

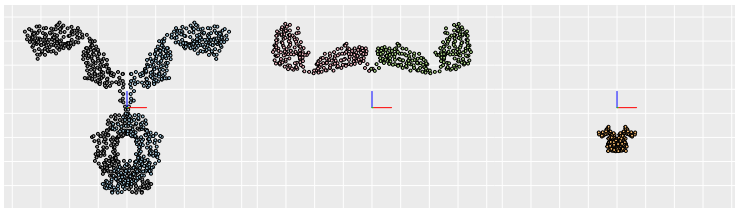
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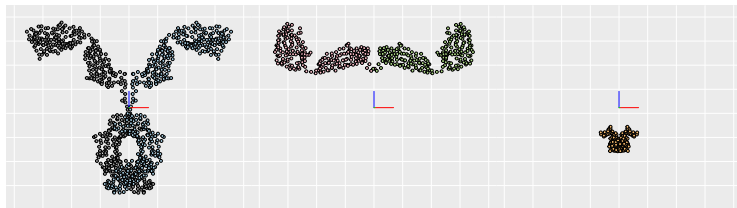
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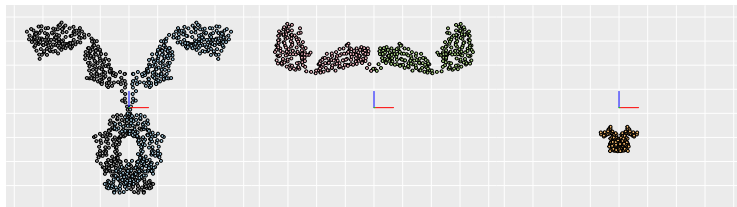


Each of these three graphical attributes is an entire subset of the data. Each is a presentation of four dimensional vectors:

$$< x(u), y(u), z(u), chainID(u) >$$

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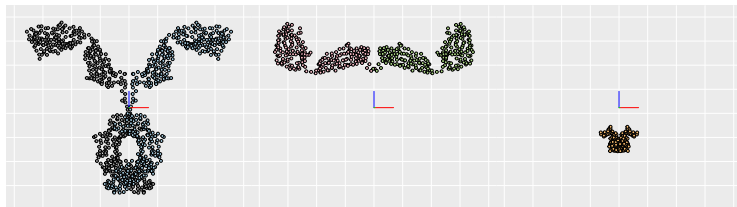
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for

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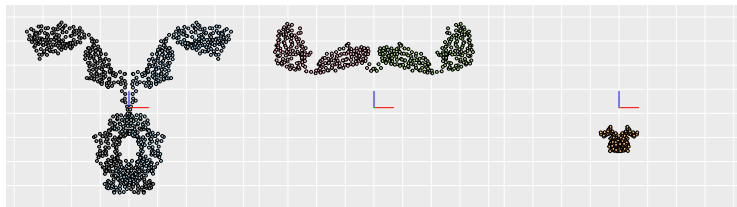
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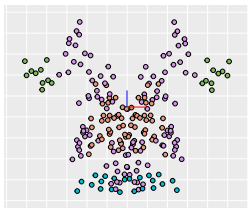
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3. $u \in \{u : u \in \mathcal{P} \text{ and } chainID(u) = "C"\}.$

Where $chainID(u)$ values are encoded by colour.

Population attributes – graphical

Or possibly zoom in on the carbohydrate chain coloured by residue:

```
p3d["active"] <- carbs  
l_scaletto_active(p3d)  
p3d["color"] <- igg1$residue  
p3d["size"] <- 10  
plot(p3d)
```



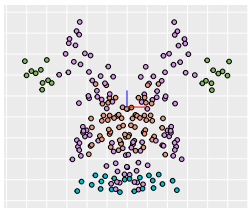
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In either case, an attribute is a summary of \mathcal{P} and as such it will always be of interest to examine how well it does and does not describe all of the units it targets in its summary.

Quick numerical attributes

Some simple attributes are easily had (and are worth checking as a habit):

```
summary(igg1)
```

```
##      recordType      name      residue      chainID residueSequenceNum
##  ATOM :1336      CA      :1336      SER      :178      C:220      Min.      : 1.0
##  HETATM: 220      C1      : 18      VAL      :122      H:452      1st Qu.: 85.0
##                                     C2      : 18      NAG      :112      I:452      Median :279.5
##                                     C3      : 18      THR      :106      L:216      Mean  :301.2
##                                     C4      : 18      PRO      :102      M:216      3rd Qu.:522.0
##                                     C5      : 18      GLY      : 98      Max.   :716.0
##                                     (Other): 130      (Other):838
##
##              x              y              z
##  Min.      : -71.18000      Min.      : -65.93      Min.      : -27.45500
##  1st Qu.: -17.32575      1st Qu.: -23.17      1st Qu.: -9.69500
##  Median : -0.01650      Median : 35.71      Median : 0.01050
##  Mean   : -0.00268      Mean   : 16.56      Mean   : 0.00856
##  3rd Qu.: 17.30550      3rd Qu.: 52.65      3rd Qu.: 9.68825
##  Max.   : 71.20500      Max.   : 75.38      Max.   : 27.52100
##
##              residueName              group
##  Serine              :178      Acidic              :110
##  Valine              :122      Basic              :146
##  N-acetylglucosamine:112      Non-polar (hydrophobic):498
##  Threonine           :106      Polar (uncharged)      :582
##  Proline             :102      Sugar              :220
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```

Each variate is given its own two columns of name : value pairs.

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##  N-acetylglucosamine:112      Non-polar (hydrophobic):498
##  Threonine    :106      Polar (uncharged)      :582
##  Proline      :102      Sugar      :220
##  Glycine      : 98
##  (Other)      :838
```

Each variate is given its own two columns of name : value pairs.

- Categorical variates show counts of values.

Quick numerical attributes

Some simple attributes are easily had (and are worth checking as a habit):

```
summary(igg1)
```

```
##      recordType      name      residue      chainID residueSequenceNum
##  ATOM :1336      CA      :1336      SER      :178      C:220      Min.      : 1.0
##  HETATM: 220      C1      : 18      VAL      :122      H:452      1st Qu.: 85.0
##                                     C2      : 18      NAG      :112      I:452      Median :279.5
##                                     C3      : 18      THR      :106      L:216      Mean  :301.2
##                                     C4      : 18      PRO      :102      M:216      3rd Qu.:522.0
##                                     C5      : 18      GLY      : 98      Max.   :716.0
##                                     (Other): 130      (Other):838
##
##              x              y              z
##  Min.      : -71.18000      Min.      : -65.93      Min.      : -27.45500
##  1st Qu.: -17.32575      1st Qu.: -23.17      1st Qu.: -9.69500
##  Median : -0.01650      Median : 35.71      Median : 0.01050
##  Mean   : -0.00268      Mean   : 16.56      Mean   : 0.00856
##  3rd Qu.: 17.30550      3rd Qu.: 52.65      3rd Qu.: 9.68825
##  Max.   : 71.20500      Max.   : 75.38      Max.   : 27.52100
##
##              residueName              group
##  Serine      :178      Acidic      :110
##  Valine      :122      Basic      :146
##  N-acetylglucosamine:112      Non-polar (hydrophobic):498
##  Threonine   :106      Polar (uncharged) :582
##  Proline     :102      Sugar      :220
##  Glycine     : 98
##  (Other)     :838
```

Each variate is given its own two columns of name : value pairs.

- ▶ Categorical variates show counts of values.
- ▶ Numeric variates show traditional summary statistics of that variate's values.

Numerical attributes

What can we learn about the distribution of the values of these variates from these numbers?

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Exercise: consider what happens to each of these measures when any variate y is transformed to $z = ay + b$ for two non-zero constants a and b .

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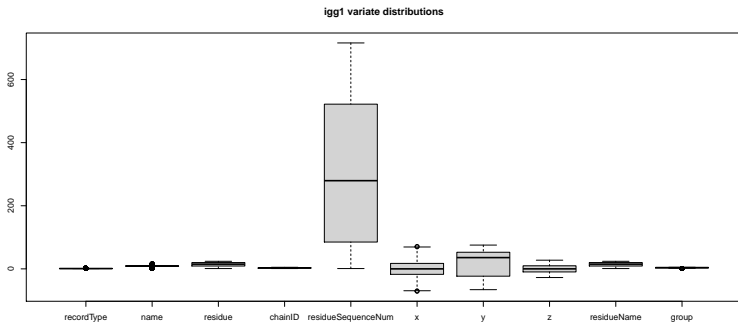
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```
boxplot(igg1, main = "igg1 variate distributions", col = "lightgrey")
```

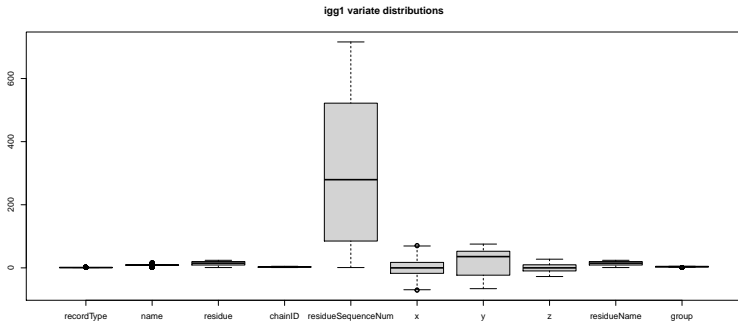


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Which is not that informative for most of the variates since they are categorical and boxplots are designed for continuous variates. Nevertheless, like `summary()` it gives a quick sense of the variates and the extent of their values.

There are other displays better suited to categorical variates.

Graphical attributes for categorical variates

Similarly, we might look at graphical attributes to summarize the distribution of values for each categorical variate.

Graphical attributes for categorical variates

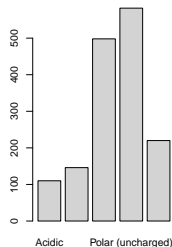
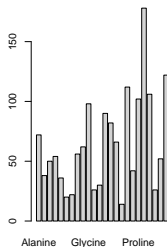
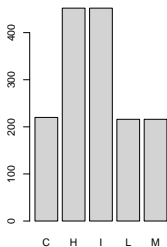
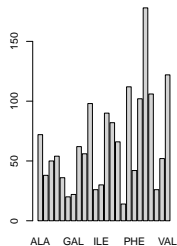
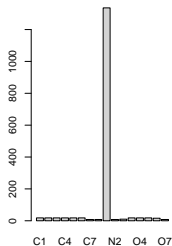
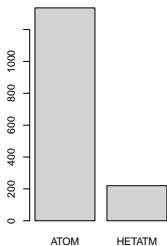
Similarly, we might look at graphical attributes to summarize the distribution of values for each categorical variate.

A bar plot for each:

```
isCatVar <- sapply(names(igg1), FUN = function(name) is.factor(igg1[,name]))
catVars <- names(igg1)[isCatVar]
nrows <- floor(sqrt(length(catVars)))
ncols <- ceiling(sqrt(length(catVars)))
savePar <- par(mfrow = c(nrows, ncols))
for (var in catVars) {
  counts <- summary(igg1[,var])
  vals <- levels(igg1[,var])
  barplot(counts, names.arg = vals, col="lightgrey")
}
par(savePar)
```

Graphical attributes for categorical variates

A bar plot for each:



Interactive graphical attributes for categorical variates

For exploratory work, it would be better if these were interactive.

```
isCatVar <- sapply(names(igg1), FUN = function(name) is.factor(igg1[,name]))
catVars <- names(igg1)[isCatVar]
# Could simply have each plot in a separate window
# or in a single window as shown here
nrows <- floor(sqrt(length(catVars)))
ncols <- ceiling(sqrt(length(catVars)))

barplotWindow <- tkoplevel() # THE WINDOW
row <- 0
col <- 0
for (var in catVars) {
  barplot <- l_hist(igg1[,var],
                    linkingGroup = "igg1",
                    title = var,
                    parent = barplotWindow)

  if (col >= ncols){
    row <- row + 1
    col <- 0
  }
  tkgid(barplot, row = row, column = col, sticky = "nesw")
  col <- col + 1}

# Configure columns to resize with window
for (col in 0:(ncols-1)){tkgrid.columnconfigure(barplotWindow, col, weight = 1)}
# Configure rows to resize with window
for (row in 0:(nrows-1)){tkgrid.rowconfigure(barplotWindow, row, weight = 1)}
# Add a title
tktitle(barplotWindow) <- "Counts for factors"
```

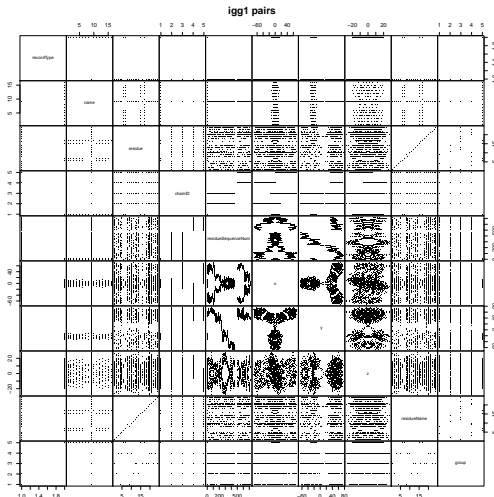
Quick graphical attributes - two dimensional

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```
plot(igg1, gap = 0, pch = ".", col = "black", main = "igg1 pairs")
```



Might be better to restrict consideration just to those variates that are not factors.

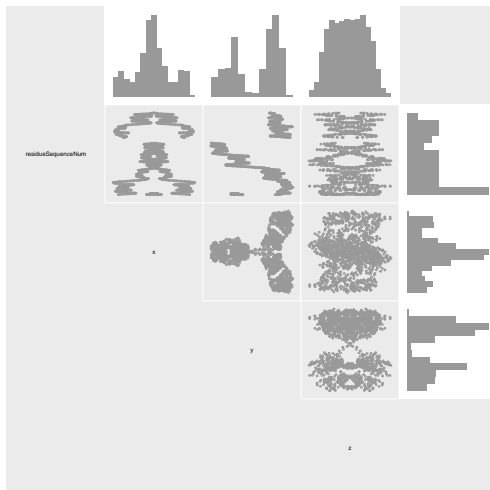
Quick graphical attributes - two dimensional interactive

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Quick graphical attributes - two dimensional interactive

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```
isCtsVar <- sapply(names(igg1), FUN = function(name) !is.factor(igg1[,name]))
ctsVars <- names(igg1)[isCtsVar]
pp <- l_pairs(igg1[,ctsVars], glyph = "ocircle", size = 1,
             showHistograms = TRUE,
             linkingGroup = "igg1", title = "Continuous pairs")
plot(pp)
```



Problem - Visible minorities in Canada 2006

Recall the minority data from loon.data.

Questions:

- ▶ What are the units u ?

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